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CERTIFICATE

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I hereby certify that annexed is a true copy of the Provisional Specification as filed on 2 November 2000 with an application for Letters Patent number 507961 made by ROBERT BARTLET ELLIOTT in trust for DIATRANZ LIMITED;DIATRANZ LIMITED;RICCARDO CALAFIORE;GUISSEPPE BASTA.

Dated 1 February 2001.

A handwritten signature in black ink that reads "Neville Harris".

Neville Harris
Commissioner of Patents



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507961

NEW ZEALAND

PATENTS ACT, 1953

PROVISIONAL SPECIFICATION

Preparation and Xenotransplantation of Porcine Islets

I, PROFESSOR ROBERT BARTLET ELLIOT an Australian citizen of 19 Laureston Avenue, Papatoetoe, Auckland, New Zealand in trust for DIATRANZ LIMITED a company duly incorporated by the laws of New Zealand, of 19 Laureston Avenue, Papatoetoe, Auckland, New Zealand, do hereby declare this invention to be described

Intellectual Property
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Introduction

The present invention relates to improvements in and/or relating to the treatment of diabetes using xenotransplantation. More particularly but not exclusively the present invention relates to the preparation of viable xenotransplantable porcine islets and/or the treatment of a mammalian patient (including humans) suffering from diabetes involving the transplantation into the mammal of viable porcine islets capable of producing insulin within the host.

Background

Type 1 (insulin-dependent) diabetes mellitus is a common endocrine disorder that results in substantial morbidity and mortality, and leads to considerable financial costs to individual patients and healthcare systems.

Treatment with insulin, while life-saving, often does not provide sufficient control of blood glucose to prevent the feared complications of the disease, which has provided the impetus for intensive research into better methods of sustaining normoglycaemia.

Among the newer treatment strategies that have been proposed, transplantation of pancreatic β islet cells, obtained either from other humans or animals, has received the most attention worldwide. This is because transplantation can restore not only the insulin-secreting unit, but also the precise fine tuning of insulin release in response to multiple neural and humoral signals arising within and beyond the islets of Langerhans.

Human islet cell transplantation is limited by the shortage of human islet tissue. The use of pig islet cells is currently viewed as the most promising alternative since:

- (a) the supply of pig cells can be readily expanded by optimising the supply of donor animals;

- (b) pig and human insulin have close structural similarities; and
- (c) physiological glucose levels in pigs are similar to those in humans.

The rationale for this treatment approach (termed 'xenotransplantation') is that the implanted pig islets have the potential to mimic the normal physiological insulin response in type 1 diabetics such that near-normal blood glucose levels may be achievable without insulin administration or with a reduced requirement for it. As a consequence, long-term diabetes complications may be prevented and patients should experience less hypoglycaemia than they do with the currently recommended 'intensive' insulin regimens.

Object

It is an object of the present invention to provide a method of preparing porcine islets which produces islets viable for xenotransplantation into a mammalian patient the islets being capable of producing insulin within a mammalian host, as well as the islet preparation so produced, or irrespectively or how produced, or a similar form.

Alternatively or additionally, it is a further object to provide a method of treating a mammalian patient suffering from diabetes which involves the xenotransplantation of porcine islets into the mammalian patient.

Alternatively or additionally, it is a further object to at least provide the public or medical community with a useful alternative approach to diabetes treatment.

Statements of Invention

In a first aspect the invention consists in a **method of preparing a xenotransplantable porcine islet preparation capable upon xenotransplantation of producing porcine insulin in an appropriate recipient mammal**, the method including or comprising:

- (i) harvesting the pancreas of piglets at or near full term gestation, and
- (ii) extracting islets from a culture of the harvested pancreas using a suitable collagenase, the culture of the harvested pancreas being (a) of mechanically reduced harvested pancreas and (b) a supportive mammalian albumin substantially free of non-human microbiological agents,

wherein the islets (at least at some stage in the performance of the method) are exposed to nicotinamide.

Preferably the piglets are at -20 to +20 days of full term gestation.

Preferably the piglets are at -7 to +10 days of full term gestation.

Preferably the mammalian albumin is human serum albumin (HSA).

Preferably the collagenase is selected from human Liberase® or porcine Liberase®.

Preferably said collagenase is human Liberase®.

Preferably the islets are treated with nicotinamide after their extraction from the pancreas.

Preferably the method includes the further step of treating the islets with IgF-1 or the N-terminal tripeptide of IgF-1 (GPE).

Preferably the pancreas and/or islets are subject to a trauma protecting agent selected from suitable anaesthetic agents.

Preferably the trauma protecting agent is lignocaine.

In another aspect the invention is **a method of preparing a xenotransplantable porcine islet preparation capable upon xenotransplantation of producing porcine insulin in an appropriate recipient mammal**, the method including or comprising the steps of:

- (I) harvesting the pancreas of piglets at or near full term gestation,
- (ii) culturing the harvested pancreas in mammalian albumin substantially free of non-human microbiological agents,
- (iii) simultaneously with step (ii) and/or after step (ii) extracting the islets using a suitable Liberase,

wherein at least one of the steps (I) to (iii) includes or is associated with the

exposure of the islets to the presence of nicotinamide.

Preferably the piglets are at -20 to +20 days of full term gestation.

Preferably the piglets are at -7 to +10 days of full term gestation.

Preferably the mammalian albumin is human serum albumin (HSA).

Preferably the collagenase is selected from human Liberase® or porcine Liberase®.

Preferably said collagenase is human Liberase®.

Preferably the islets are treated with nicotinamide after their extraction from the pancreas.

Preferably the method includes the further step of treating the islets with IgF-1 or the N-terminal tripeptide of IgF-1 (GPE).

Preferably the pancreas and/or islets are subject to a trauma protecting agent selected from suitable anaesthetic agents.

Preferably the trauma protecting agent is Lignocaine.

In another aspect the invention consists in a **method of preparing a xenotransplantable porcine islet preparation capable upon xenotransplantation of producing porcine insulin in an appropriate recipient mammal**, said method including all comprising:

- (I) harvesting the pancreas of piglets at -20 - +20 days of full term gestation
- (ii) mechanically reducing the harvested pancreas in the presence of an islet trauma protecting agent and producing a culture of the mechanically reduced harvested pancreas supported by human serum albumin (HSA),
- (iii) isolating the islet cells from the culture of mechanically reduced harvested pancreas using a Liberase selected from human Liberase or porcine Liberase,
and
- (iv) encapsulating the islet cells with a biocompatible xenotransplantable material, said material *in vivo* being both glucose and insulin porous, wherein nicotinamide is introduced to the islets or islet cells prior to encapsulation at any one or more stages of the procedure, and
wherein, at least to the extent required to convert porcine non-insulin producing

islet cells to porcine insulin producing islet cells, at some stage in the procedure presenting the islets and/or islet cells to IgF-1 or the N-terminal tripeptide of IgF-1 (GPE).

Preferably an antibiotic is associated with the islet cells.

Preferably said antibiotic is ciproxin.

Preferably said islet trauma protecting agent is lignocaine.

Preferably said biocompatible material is a suitable alginate.

Preferably the alginate is in ultra pure form.

Preferably each islet or grouping of islets is entrapped in an *in vivo* insulin and glucose porous biocompatible alginate or alginate-like surround.

Preferably the encapsulation provides a surround which prevents, once implanted, direct tissue contact with the islets.

Preferably each encapsulation involves presenting islets and a suitable alginate solution into a source of compatible cations thereby to entrap the islets in a cation - alginate gel.

Preferably said cation alginate gel is calcium-alginate gel.

Preferably said alginate used in the solution is sodium alginate, and the islets and sodium alginate solution is presented as a droplet into a bath of suitable cations.

Preferably the islets and sodium alginate solution is of 1.6% w/w.

Preferably the islets and sodium alginate solution is presented as a droplet through a droplet generating needle.

Preferably the suitable cations are calcium chloride.

Preferably the gel encased islets are coated with a positively charged material and thereafter are provided with an outer coat of a suitable alginate.

Preferably the positive charging material is poly-L-ornithine.

Preferably the gel entrapping the islets within the outer coating is then liquified.

Preferably the liquification involves or comes about by the addition of sodium citrate.

Preferably the encapsulation produces capsules.

Preferably the capsules contain a plurality of islet cells.

Preferably the capsules contain substantially three islet cells.

Preferably the capsules have a diameter of substantially from about 300 to 400 microns.

Preferably following liquification of the alginate entrapping the islets there are the further steps of:

- washing the capsules

- further coating the capsules with alginate to neutralize any residual change on the poly-L-ornithine coating and prevents direct contact of the poly-L-ornithine with tissues when the entire capsule is transplanted.

Preferably the alginate has been produced via a process involving the steps of:

Seaweed harvest → Washing → Alginate extraction → Filtration
→ Precipitation → Drying.

Preferably the alginate is Keltone LVCR (produced by I SP Alginates Inc US or similar) and has the following specifications:

1. Viscosity: 2% - 100-300 cps (Brookfield 25°C, speed 3,60 rpm)
2. pH: 6.4-8.0
3. Chemical analysis:

total heavy metals <40 ppm

Pb: <10 mg/kg (ppm)

As: <1.5 mg/kg (ppm)

4. Endotoxin level - measured by LAL test (at University of Perugia): 39 EU/g

[NB. Any level below 100 EU/g in this test is considered endotoxin-free].

5. Microbiology Limits

Bacteria <200 cfu/g Salmonella absent in 25g

Yeast <100 cfu/g Staphylococcus auseus absent in 1.0g

Mold <100 cfu/g Pseudomonas aeruginosa absent in 1.0g

Coliform neg by MPN

E.Coli absent in 25g

6. Mannuronic acid (M) content: M fraction (F_m) 70%
7. Guluronic acid (G) content: G fraction (F_G) 30%
8. Quality System: Keltone LVCR is manufactured according to ISO 9002.

In another aspect the present invention is a **xenotransplantable preparation** being or including viable porcine islets prepared according to a method of the present invention.

In yet another aspect the invention is a **xenotransplantable capsule** having viable islet cells from a -20 to +20 full term gestation piglet capable of producing porcine insulin in response to glucose within a recipient mammal within its biocompatible encapsulating material or materials, the capsule being such that the encapsulation is such as to prevent tissue contact with said islet cell(s) by tissue of a recipient mammal yet *in vivo* will allow glucose entry to the islet cells and the egress of porcine insulin from such islet cells, the islet cells having been isolated from a culture supported by HSA using a suitable Liberase® and prior to encapsulation having been exposed to nicotinamide.

Preferably said culture has been of mechanically reduced harvested pancreatic tissue, such tissue having been exposed to a trauma reducing agent.

Preferably said encapsulation has been of islet cells in the presence of a suitable antibiotic.

In yet another aspect the invention is a **method of treating a mammalian patient predisposed to or suffering from diabetes** which involves the xenotransplantation into such patient at least one capsule of the present invention.

In yet another aspect the invention is a **method for the treatment of a mammalian patient suffering from or predisposed to diabetes**, said method including or comprising the steps of:

- (A) (i) harvesting the pancreas of piglets at or near full term gestation,
- (ii) culturing the harvested pancreas in Mammalian Albumin

substantially free of non-human microbiological agents,

(iii) simultaneously with step (ii) and/or after step (ii), extracting the islets from the harvested pancreas using a suitable Liberase, wherein the islets (at least at some stage in the performance of (A)) are exposed to nicotinamide;

(B) (i) encapsulating the islets prepared by (A) with a suitable encapsulation material that allows both glucose and insulin movement therethrough, and

(ii) implanting the encapsulated porcine islets into the recipient mammal.

Preferably there is included also the step of administering nicotinamide to the recipient mammal prior to or after the implantation step.

Preferably the method further includes the step of prescribing for the patient, prior to or after the implantation step, a casein-free diet (as described herein).

Preferably the method further includes the step of subjecting the patient prior to or after the implantation step to a cholesterol lower drug regime.

Preferably the cholesterol lowering drug is of the "statin" family

Preferably said cholesterol lowering drug is pravastatin or simvastatin.

In another aspect the invention is part or all of a procedure of Figure 1.

Definitions

As used herein:

- "*Administering*" includes self-administering;
- "*Casein-free*" when referring to milk as used herein refers to milk which does not contain a diabetogenic factor, particularly to milk containing no variant of β -casein which stimulates diabetogenic activity in humans. With reference to International PCT Application WO 96/14577, a non-diabetogenic variant for example, may be the A2 variant of β -casein. The full contents of PCT/NZ95/00114 (WO 96/14577) and PCT/NZ96/00039 (WO 96/36239) are here included by way of reference.

“Casein-free” as used herein in respect of dietary considerations means at least a substantial avoidance (preferably total avoidance) of such milk containing or derived diabetogenic factors.

- The *N-terminal tripeptide of Ig F-1* or “GPE” is a tripeptide (gly-pro-glu) derived from Ig F-1.
- “*mammalian albumin*” as used herein means serum albumin from mammals, preferably human serum albumin (HSA).
- “*appropriate collagenase*” means preferably Liberase®, ideally human or porcine, ideally Liberase H®.
- “*mechanically reduced*” as used herein includes any process where pancreatic tissue is increased in surface area eg, mechanical or water jet shredding, grinding, mincing, etc...

1. General

The present invention recognises the ability to source appropriate islets from piglets which have similar structural similarities of insulin to humans, and similar physiological glucose levels to humans. The piglets used are at or near full term gestation. The islets are converted into an appropriate xenotransplantable source of islets with viability in a human being by following certain procedures in respect of the harvesting and extraction of the islets, the treatment of the islets prior to xenotransplantation as well as regimes of use of such islets.

The major advantage of porcine islet cell transplantation over human islet cell transplantation is that the islet cell source can be readily expanded, and the biosafety of the cells can be thoroughly explored prior to transplantation. From a practical viewpoint, pancreas removal and islet cell isolation can be performed expeditiously in an ideal environment.

Important considerations relevant to the use of porcine islet cells in transplantation approaches for type 1 diabetes include the following:

- The structural and biological similarities of porcine and human insulin
- The fact that porcine insulin has been used to treat diabetes for several decades (and has only been replaced by human sequence insulin relatively recently); and
- The similarity of physiological glucose levels in pigs and humans. (Weir & Bonner-Weir 1997). This effectively means that pig islet cells can be expected to react similarly to their human counterparts in maintaining equivalent blood glucose concentrations.

2. The Nature of the Disease causing Diabetes

Successful long-term allotransplantation of human islets can be achieved in over 80% of patients when the disease is caused by non-immune processes. In contrast, even islets obtained from a non-diabetic twin cannot reverse autoimmune diabetes long-term in the diabetic twin member. This emphasises the critical role of autoimmunity in the failure of islet transplantation. This observation has been validated in allotransplantation of rodents with diabetes caused by autoimmunity as compared with diabetes due to pancreatectomy or chemical β cell destruction. No large animal model of autoimmune diabetes exists. It is possible that the use of islets from different species (xenotransplantation) could avoid autoimmune destruction of transplanted islets, as the immune process of xenotransplant rejection is different to that of allotransplant rejection, but this is entirely hypothetical in humans.

3. Isolation and Preparation of Porcine Islet Cells for Xenotransplantation

3a. Animal Source and Transportation

All animals intended as a source of pancreatic tissue for xenotransplantation are obtained from a specific pathogen-free (SPF) pig breeding facility which is maintained in accordance with the American Association for Accreditation of Laboratory Animal Care (AAALAC). The facility

maintains a high-health status colony with excellent standards of husbandry, and operates a record system that is readily accessible and archived indefinitely. Donor sows and sires are selected with the underlying objective of producing strong heterosis in donor litters.

3b. Isolation and Purification of Islet Cells

Following surgical removal, the donor pancreases are transferred to a cleanroom facility for further processing in a cold plastic container in 50ml tubes containing cold Hanks' Balanced Salt Solution (HBSS) with 0.2% human serum

albumin (HSA) added. Blood samples from each donor are sent for virology testing and toxoplasma serology. Samples from each organ are kept in a freezer at -80°C for future testing if necessary.

3c. Digestion

The islet cells are isolated by standard collagenase digestion of the minced pancreas via the procedure documented by Ricordi et al. (1990), though with some modifications. Using aseptic technique, the glands are distended with Liberase® (1.5mg/ml), trimmed of excess fat, blood vessels and connective tissue, minced, and digested at 37°C in a shaking water bath for 15 minutes at 120 rpm. The digestion is achieved using lignocaine mixed with the Liberase® solution to avoid cell damage during digestion. Following the digestion process, the cells are passed through a sterile 400mm mesh into a sterile beaker. A second digestion process is used for any undigested tissue.

We have determined that much greater yields per neonatal pig pancreas can be obtained using either pig or human Liberase™ (eg; sourced in New Zealand from Roche) rather than collagenase. Whilst there is disclosure in "*Improved Pig Islet Yield and Post-Culture Recovery Using Liberase P1 Purified Enzyme Blend*", T J Cavanagh et al. Transplantation Proceedings 30, 367 (1998) and in "*Significant Progress In Porcine Islets Mass Isolation Utilizing Liberase® HI For Enzymatic Low-Temperature Pancreas Digestion*", H. Brandhorst et al.

Transplantation Vol 68, 355-361 No. 3, August 15, 1999 the yields therefore therein are low compared to those we have discovered. If, for example, in following the procedure of Brandhorst et al. there is a yield increase of islets over collagenase of from 400 to say 800 with the procedure using human Liberase ® (ie; Liberase ® HI) as in the Brandhorst et al. procedure but confined to neonatal porcine islets such as those as 7 days post delivery extra ordinarily larger yields are possible, namely, the equivalent to from 400 which would be the case with crude collagenase to 30000 which as can be seen as very much greater than that to be expected from following the procedure of Brandhorst et al. with pigs.

3d. Washing and Culture

The digested tissue is washed three times, and seeded into cell culture media RPMI 1640 to which is added 2% human serum albumin (HSA), 10 mmol/L nicotinamide, and antibiotic (Ciproxin).

3e. Quality Control Procedures

To exclude any contamination of the tissue, quality control procedures are undertaken on cell culture samples after isolation and before encapsulation (further details are given in SOP P101). Three days after isolation, the cell culture is tested for microbiological contamination by accredited laboratories. Testing for porcine endogenous retrovirus (PERV) is undertaken at the Virology Laboratory, Auckland Hospital.

The *islet yield* is determined via dithizone (DTZ) staining of the cells, as specified in SOP Q200. Dithizone is a zinc-chelating agent and a supravital stain that selectively stains zinc in the islets of Langherhans, producing a distinctive red appearance.

The *viability* of the islet cells is determined using acridin orange and propidium iodide, as specified in SOP Q201. Acridin orange is a fluorescent stain that readily passes through all cell

membranes to stain the cytoplasm and nucleus. Bright green fluorescence in both the nucleus and cytoplasm on exposure to ultraviolet (UV) light denotes intact live cells. Conversely, propidium iodide is a fluorescent stain that cannot pass through an intact membrane. It emits a bright red fluorescence when exposed to UV light, and the presence of propidium iodide in a cell nucleus indicates severe damage or a dead cell.

3f. Determination of *in vitro* Insulin Secretory Capacity

Static glucose stimulation (SGS) is used to assess *in vitro* function of the porcine islets by exposing them to low and high concentrations of glucose and theophylline. Determination of the *in vitro* insulin secretory capacity is undertaken on both free islets (after 3 days in culture) and after their subsequent encapsulation or confinement.

4. Xenotransplantation

4a. The viability of the islets for xenotransplantation

The processes by which islets are purified prior to transplantation are traumatic to these highly specialised tissues. Such trauma can induce necrosis or apoptosis – the latter being quite delayed.

Further trauma may result from encapsulation. Processes used by us in both the preparation of islets and their encapsulation have been optimised to ensure minimal damage to the islets. Such procedures have ensured zero warm ischaemia (compared with hours with most human islet preparations), have involved the use of nicotinamide to enhance successful *in vitro* explanation, have involved minimal incubation time with collagenase or liberase, have involved swift non-traumatic encapsulation technology, have involved the use of IgF-1 (or the GPE tripeptide thereof), the use of an anaesthetic such as lignocaine, and the use of an antibiotic such as ciproproxin etc.

Our preferred preparation preferably uses neonatal (7-day old) islets which is crucial in both limiting islet trauma during purification, and assuring sufficient maturation of the islets for stimulated insulin production.

The IgF-1 (Human Insulin-like Growth Factor I) is used in order to induce unmatured porcine islets to mature to their insulin-producing form. IgF-1 is a potent mitogenic growth factor that mediates the growth promoting activities of growth hormone postnatally. Both IgF-1 and IgF-2 are expressed in many cell types and may have endocrine, autocrine and paracrine functions. The preferred form of IgF-1 we have found to be the amino-terminal tripeptide glycine-proline-glutamate of IgF-1 (GPE).

4b. Alginate Encapsulation Procedure

Sodium alginate used for this procedure is extracted from raw material sources (seaweed) and prepared in a powdered ultrapure form. The sterile sodium alginate solution (1.6%) is then utilised at the Diatranz Islet Transplant Centre to manufacture encapsulated islets. The encapsulation procedure (University of Perugia) involves extruding a mixture of islets and sodium alginate solution (1.6%) through a droplet generating needle into a bath of gelling cations (calcium chloride). The islets entrapped in the calcium-alginate gel are then coated with positively charged poly-L-ornithine followed by an outer coat of alginate (0.05%). The central core of alginate is then liquefied by the addition of sodium citrate. Most capsules contain 3 islet cells and have a diameter of 300 to 400 μ m.

The encapsulated islets are kept in cell culture, and then checked for contamination, insulin release and viability before transplantation. They are only released for transplantation if all quality control tests are negative.

4c. Drugs used in the recipient

Transplantation does not require and avoids the need for cytotoxic agents to suppress the immune system. Such agents are able to enter the alginate microcapsule and cause islet toxicity, as well as causing systemic toxicity. Instead, nicotinamide and a special diet are used (for rationale, see section 1.4 below).

The transplantation procedures of our earlier patent specification have the ability over a period prior to rejection of providing porcine insulin. In this respect, we ourselves conducted clinical trials.

Four type 1 diabetic adolescents received 10,000 free islets/kg bodyweight by intraperitoneal injection. The islets were located from term piglets using the standard collagenase digestion, purification and culture techniques described in section 3.2. All four recipients received oral nicotinamide (1.5 g/day) and a casein-free as herein defined diet both pre- and post-transplantation. A prompt reduction in insulin requirements, which was not clearly dose-related, was noted in the first week after transplantation. The reduction in insulin dosage range from 21 to 32%, and the response lasted for up to 14 weeks. However, insulin doses subsequently returned to their previous levels.

The most likely reason for the transplant failure in these patients was chronic rejection. However, no adverse effects were noted.

We have now shown alginate-encapsulated porcine islet cell transplants in two human diabetic patients, prolonged functioning of the transplants. The islets were transplanted by intraperitoneal injection, one patient receiving 15,000 IEQ/kg (total 1,300,000 islets) and the other 10,000 IEQ/kg (total 930,000 islets). Both patients were treated pre- and post-transplantation with oral nicotinamide and a soy-based/casein-free as herein defined diet.

The preferred procedure as shown in Figure 1 was used for the preparation, the encapsulation being as aforesaid. Islet cells of -7 days to +10 days full gestation were used.

Description of the Drawings

This invention may also be said broadly to consist in the parts, elements and features referred to or indicated in the specification of the application, individually or collectively, and any or all combinations of any two or more of said parts, elements or features, and where specific integers are mentioned herein which have known equivalents in the art to which this invention relates, such known equivalents are deemed to be incorporated herein as if individually set forth.

The invention consists in the foregoing and also envisages constructions of which the following gives examples.

Preferred forms of the present invention or examples of working will now be described with reference to the accompanying drawings in which:

Figure 1 shows a preferred procedure for harvesting, isolating and preparing islet cells (with either confinement or encapsulation) and the associated treatment regime for a diabetic human patient in order to receive ongoing benefit from the xenotransplantation,

Figure 2 shows the effect of collagenase from various sources on islet yield and function,

Figure 3 shows the stimulation index of Liberase® against Collagenase clearly showing that Liberase® preparations (both human and porcine at suitable concentrations) gave higher yields and function in vitro than an optimised concentration of Collagense P,

Figure 4 shows the stimulation index of free islets when comparing the use of ciproxin against a penicillin/streptomycin mix and against a control of no antibiotics,

Figure 5 shows the results of exposure of neonatal porcine islets in culture with GPE in comparison with control cells.

5. Examples

5a. Examples of use of IgF-1

- porcine islets in culture which were exposed to IgF-1, increased their insulin response to glucose, by up to a 3-fold increase.

	Incubated 24hrs with 0.1ug/ml IgF-1 after isolation	CONTROL- no IgF-1
Insulin secretion In response to 19.4mM Glucose +10mM Theophylline After 3 days culture Post isolation	236uU/hr/100IEQ	75.2uU/hr/100IEQ

- A concentration of 0.1ug/ml IgF-1 in culture is sufficient to produce optimal insulin secretion during glucose challenge. No further benefit was achieved by increasing the concentration of IgF-1.

	Incubated 24hrs with 0.1ug/ml IgF-1	Incubated 24hrs with 1.0 ug/ml IgF-1
Insulin secretion In response to 19.4mM Glucose +10mM Theophylline After 3 days culture Post isolation	58uU/hr/100IEQ	56.8uU/hr/100IEQ

- Variations on the duration of IgF-1 exposure were tried on the porcine islet cells. However no increased benefit was found on culturing the islets with IgF-1 beyond a 24hrs period, post isolation.

	Incubated 7 days With 0.1ug/ml IgF-1	Incubated 24hrs with 1.0 ug/ml IgF-1
Insulin secretion In response to 19.4mM Glucose +10mM Theophylline 7days post isolation	58uU/hr/100IEQ	57.5uU/hr/100IEQ

- This increased insulin production persisted to 14 days post IgF-1 exposure. Longer durations are yet to be investigated.

	14 days post IgF-1 Exposure	3 days post IgF-1 Exposure
Insulin secretion In response to 19.4mM Glucose + 10mM Theophylline	1.3-fold increase Compared to control	1.5-fold increase Compared to control

- Withdrawal of Nicotinamide from the culture media eliminated the benefit of IgF-1 on islet insulin production.

	Incubated 3 days With 0.1ug/ml IgF-1 Without Nicotinamide	Incubated 3 days With culture Media Without Nicotinamide
Insulin secretion In response to 19.4mM Glucose +10mM Theophylline After 3 days culture Post isolation	47.6uU/hr/100IEQ	55.9uU/hr/100IEQ

- A concentration of 0.1ug/ml IgF-2 during culturing appeared to increase insulin production of porcine islet cells, after an initial exposure of 24 hrs. However, this increase was transient to 3 days post exposure.

	Incubated 24hrs With 0.1ug/ml IgF-2 day 1.	Control
Insulin secretion In response to 19.4mM Glucose +10mM TheophyllineAfter 3 days culture Post isolation	105.8/100IEQ	75.2r/100IEQ

	Incubated 24hrs With 0.1ug/ml IgF-2 day 1.	Control
Insulin secretion In response to 19.4mM Glucose +10mM TheophylineAfter 3 days culture Post isolation	32uU/hr/100IEQ	39.8 uU/hr/100IEQ

- Prolonged exposure to IgF-2 beyond 24hrs, failed to increase the insulin production of the islet cells in response to glucose.

	Incubated 24hrs With 0.1ug/ml IgF-2 day 1.	Control
Insulin secretion In response to 19.4mM Glucose +10mM TheophylineAfter 3 days culture Post isolation	105.8/100IEQ	75.2r/100IEQ

	Incubated 7 days With 0.1ug/ml IgF-2	Control
Insulin secretion In response to 19.4mM Glucose +10mM TheophylineAfter 7 days culture Post isolation	38.4uU/hr/100IEQ	39.8uU/hr/100IEQ

5b. Effect of N-Terminal Tripeptide (GPE) of Insulin like growth factor (IGF-1) on the function of neonatal porcine islet cells.

GPE is a tripeptide (gly-pro-glu) derived from IGF-1. It is a novel neuroactive peptide with a potent effect on acetylcholine and dopamine release in cortical slices. The studies done using GPE support the concept that the proteolytic products of the IGF-1 precursor play a role in the regulation of brain functions.

The aim of this example was to present the effect of GPE on the function of isolated porcine islets in vitro.

Method

- Islet cell isolation with 2 pancreases;
- Isolation following the previously discussed protocol;
- RPMI media added with Ciproxin, nicotinamide, Human serum albumin
- IGF1- Long R 3 IGF-1 Media Grade, GroPep (AU100), stock solution 100 ug/ml (in 10 mM HCl): dilute further in RPMI medium to the final concentrations: 1ug/ml (1: 100), 0.5 ug/ml (1: 200) and 0.1 ug/ml (1:1000).
- GPE, IGF1 (1-3), Bachem AG, Lot No.0538925, stock solution of 100 ug/ml (in water): dilute further in RPMI medium to the final concentrations: 1ug/ml (1:100), 0.1 ug/ml (1: 1000) and 0.01 ug/ml (1: 10 000)
 - 1. IGF1 0.1ug/ml
 - 2. IGF1 0.5ug/ml
 - 3. IGF1 1.0ug/ml
 - 4. GPE 0.01ug/ml
 - 5. GPE 0.1ug/ml
 - 6. GPE 1.0ug/ml

Keep the cells 3 days in culture before Static Glucose Stimulation (SGS). SGS involves exposure of the cells to low and high concentration of glucose to check insulin production. Using 0.1 ug/ml concentration add IGF1 and GPE to two plates 24 hours before SGS (day 2 after isolation)

Results of Example Sb

Exposure of neonatal porcine islets in culture to GPE increased the insulin response to glucose

up to 11.5 fold compared with the control cells.(Stimulation Index control 13.3 compared to 24.8 when GPE was used) Viability of the cells was >85% DTZ, AO/PI staining)

A concentration of 0.01ug/ml of GPE in culture is sufficient to produce optimal response during glucose challenge. No further benefit was achieved by increasing the concentration of GPE in culture. See Figure 5 below.

The results suggest that GPE could be used during porcine islet cell culture to improve the quality and function of the cells before transplantation. Furthermore GPE is a novel neuroactive peptide found in human brain.

5c. Examples of the effect of lidocaine when used during porcine pancreatic digestion, on islet yield and viability.

Lidocaine is a membrane stabiliser and phospholipase A2 inhibitor. When used at a 1mM concentration during Collagenase digestion of 7d old porcine pancreas, a 2-fold increase in islet yield is produced.

Islet endocrine function was assessed after 3 days in culture via static glucose stimulation. Islets isolated with Lidocaine during digestion produced a 3-fold increase in insulin secretion in response to glucose challenge.

	Collagenase alone	Collagenase + 1mM Lidocaine
Average islet yield	40,960 IEQ/g	88,183 IEQ/g

	Collagenase alone	Collagenase + 1mM Lidocaine
Insulin secretion in response to 19.4mM Glucose +10mM Theophylline After 3 days culture Post isolation	46.4 uU/hr/100IEQ	163.8 uU/hr/100IEQ

Conclusion: The use of Lidocaine during pancreatic digestion increases the insulin production/g of pancreas by 6-fold.

5d. Examples of the effects of Ciproxin on Islet function as assessed by static glucose stimulation.

Freshly prepared neonatal pig islets were prepared by standard isolation procedure and cultured for two days in RPMI medium with standard additions.

Streptomycin (100mg/ml)and Penicillin (100U/ml) were included in one flask and Ciproxin (3 mcg/ml) in another.

The islets were harvested and an aliquot subjected to stimulation with theophylline and high glucose.

The comparative insulin release from the islets---a measure of viability is shown in Figure 4.

5e. Examples of the effects of collagenase from various sources on islet yield and function

Pancreases of neonatal piglets aged 7 days were obtained as above and islets extracted by the same process, varying only the source and amount of collagenase. The yield/gram of pancreas is shown in the Figure.

Islets extracted using these variations in collagenase source and amount were assessed for viability using propidium iodide and dithizone for insulin content.

DTZ staining >85%

AO/PI >85%

The islets were then assessed for functionality by static glucose stimulation as above. The

results are shown in the Figure below.

It is apparent that the Liberase® preparations at suitable concentrations gave higher yields and function in vitro than the previously optimised concentration of Collagenase P.

5f. Examples of the comparative effectiveness of islets prepared with Liberase P or H in vivo

Islets prepared with the best concentration of Liberase® P and H in this way were injected intraperitoneally into CD1 mice made diabetic by intravenous streptozotocin. The dose used was 10 islets/g body weight of mouse. Ten days after such treatment the number of mice no longer diabetic was assessed.

1/7 of the mice treated with the islets isolated with Liberase® P and 4/7 of those isolated with Liberase H were non diabetic.

Similar experiments were performed using spontaneously diabetic NOD mice. Of the surviving mice at 10 days after transplantation 3/7 of the Liberase P treated islets and 3/3 of the Liberase H islets were no longer diabetic.

5g. Example of Islet Encapsulation Procedure

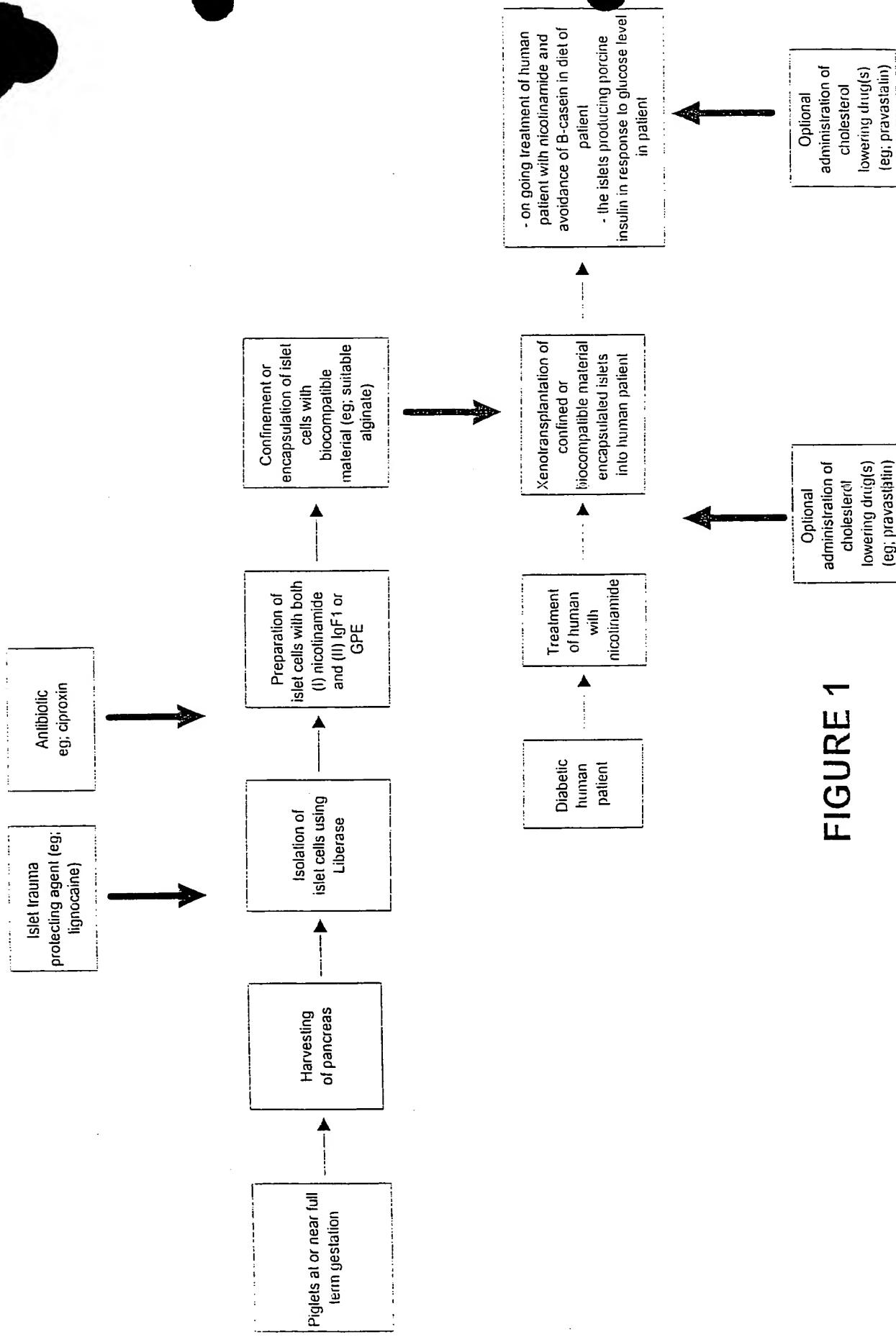
The novel medium size microcapsules (300-400 μ MSM) are prepared by atomizing the islet-alginate suspension through a special microdroplet generator.

Sodium alginate used for this procedure is extracted from raw material sources (seaweed) and prepared in powdered ultrapure form (Keltone LVCR).

The encapsulation procedure involves extruding a mixture of islets and sodium alginate solution (1.6%) through a droplet generating needle into a bath of gelling cations (calcium

chloride). The islets entrapped in the calcium-alginate gel are then coated with positively charged poly-L-ornithine followed by an outer coat of alginate (0.05%). The central core of alginate is then liquified by the addition of sodium citrate. Most capsules contain 3 islet cells and have a diameter of 300 to 400 μ m.

The encapsulated islets are kept in cell culture, and then checked for contamination, insulin release and viability before transplantation.



Porcine Islet yield Collagenase P Vs Liberase

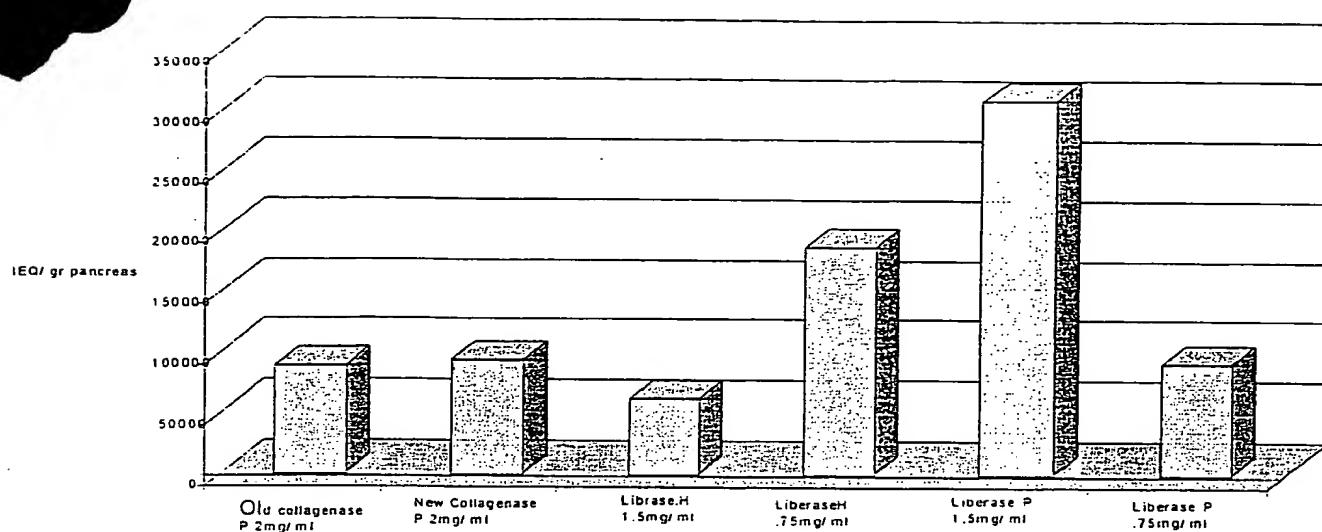


FIGURE 2

Stimulation Index Liberase Vs Collagenase

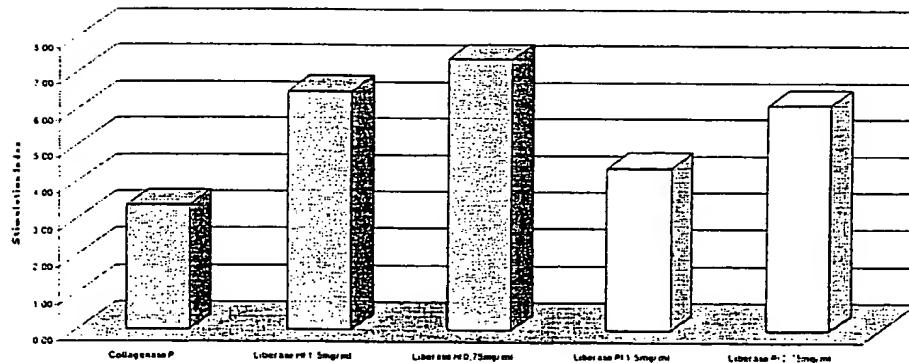
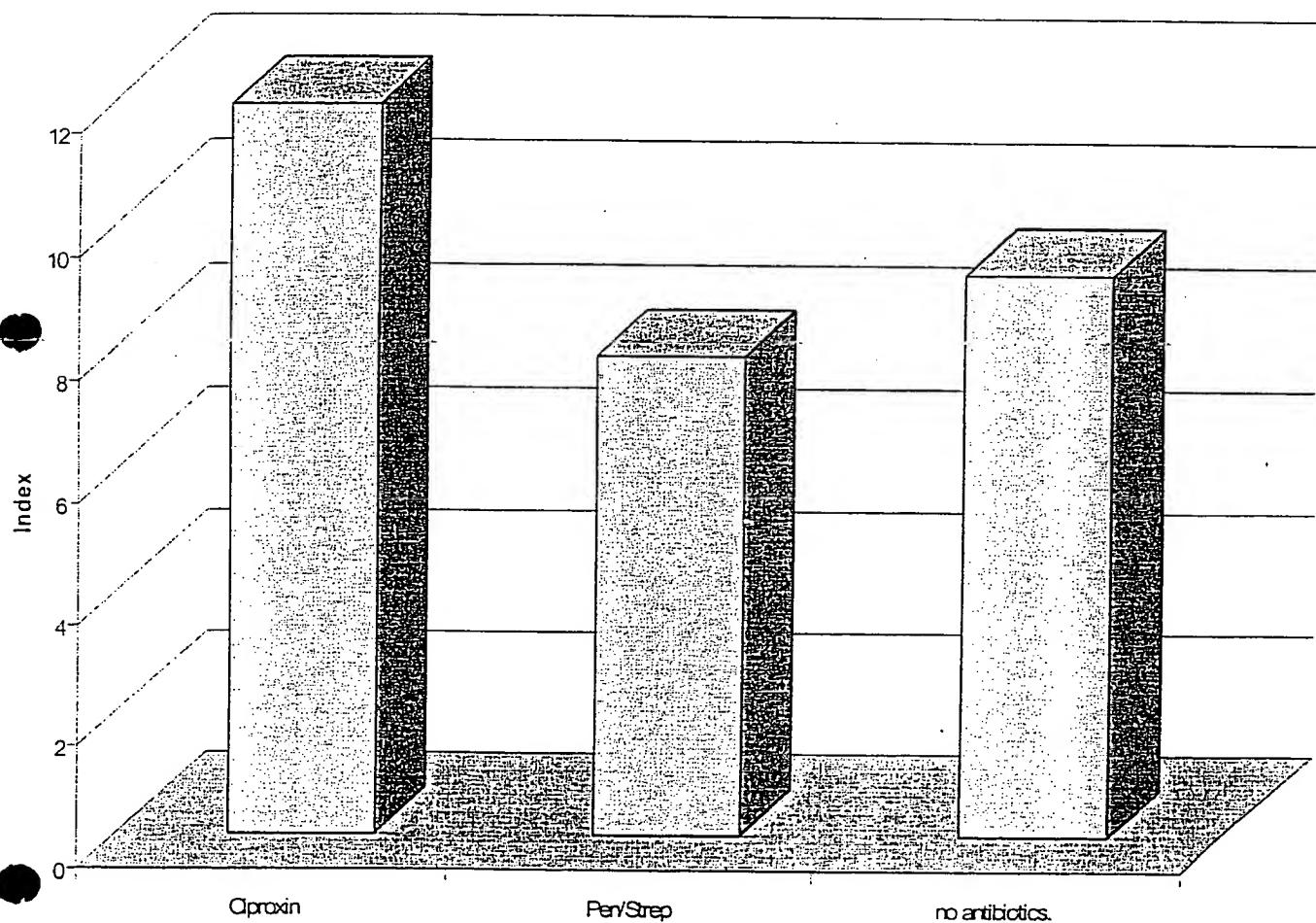


FIGURE 3

FIGURE 4

Stimulation Index Free Islets Ciproxin Study



Effect of N-Terminal Peptide (GPE) on Islet Cell Culture

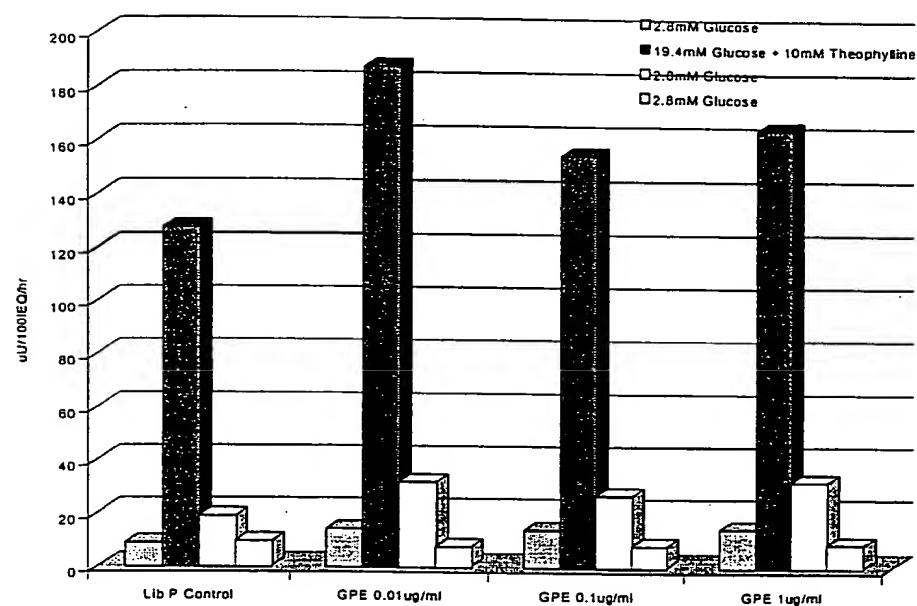


FIGURE 5